

FORM PTO1390
(REV 10-92)

U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTORNEY'S DOCKET NO.

TRANSMITTAL LETTER TO THE UNITED STATES
DESIGNATED/ELECTED OFFICE (DO/EO/US)

153-5916/PCT

09/868871

INTERNATIONAL APPLICATION NO.
PCT/IB97/00770INTERNATIONAL FILING DATE
24/06/1997 (24 June 1997)PRIORITY DATE CLAIMED
27/06/1996 (27 June 1996)

TITLE OF INVENTION

STABILIZER COMPOSITIONS

APPLICANT(S) FOR DO/EO/US

JAN MALIK ET AL.

Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items under 35 U.S.C. 371:

1. ☒ This express request to immediately begin national examination procedures (35 U.S.C. 371(f)).
2. ☒ The U.S. National Fee (35 U.S.C. 371(c)(1)) and other fees as follows:

CLAIMS	(1) FOR	(2) NUMBER FILED	(3) NUMBER EXTRA	(4) RATE	(5) CALCULATIONS
	TOTAL CLAIMS	11 - 20 =	0	X \$18.00	
	INDEPENDENT CLAIMS	1 - 3 =	0	X \$78.00	
	MULTIPLE DEPENDENT CLAIMS (if applicable)			+ 260.00	
	BASIC NATIONAL FEE (37 CFR 1.492(a) (1)-(4):				
	<input checked="" type="checkbox"/> For filing with EPO or JP search report (37 CFR 1.492(a)(5))			\$ 840.00	840.00
	<input type="checkbox"/> International Preliminary Examination fee paid to USPTO (37 CFR 1.482)			670.00	
	<input type="checkbox"/> No international preliminary examination fee paid to USPTO (37 CFR 1.482) but international search fee paid to USPTO (37 CFR 1.445(a)(2))			760.00	
	<input type="checkbox"/> Neither international preliminary examination fee paid to USPTO (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO			970.00	
	<input type="checkbox"/> International preliminary examination fee paid to USPTO (37 CFR 1.482) and all claims satisfied provisions of PCT Articles 33(2)-33(4)			96.00	
	Surcharge of \$130.00 for furnishing the National fee or oath or declaration later than <input type="checkbox"/> 20 <input checked="" type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(e)).				
	TOTAL OF ABOVE CALCULATIONS				= 840.00
	SUBTOTAL				+ 840.00
	Processing fee of \$130.00 for furnishing the English translation later than <input type="checkbox"/> 20 <input type="checkbox"/> 30 months from the earliest claimed priority date (37 CFR 1.492(f)).				
	TOTAL NATIONAL FEE				\$840.00
	Fee for recording the enclosed assignment (37 CFR 1.21(h)).				+ 40.00
	TOTAL FEES ENCLOSED				\$880.00

- a. ☐ A check in the amount of \$_____ to cover the above fees is enclosed.
- b. ☒ Please charge my Deposit Account No. 03-2060 in the amount of \$ 880.00 to cover the above fees. A duplicate copy of this sheet is enclosed.
- c. ☒ The Commissioner is authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 03-2060. A duplicate copy of this sheet is enclosed.

0936371 00001 105090 14339360

09/868871

3. A copy of the International Application as filed (35 USC 371(c)(2))
- a. ☒ is transmitted herewith.
 - b. ☐ is not required, as the application was filed in the US Receiving Office (RO/US).
 - c. ☐ has been transmitted by the International Bureau.
4. ☐ A translation of the International Application into English (35 USC 371(c)(2)).
5. ☐ Amendments to the claims of the International Application under PCT Article 19 (35 USC 371(c)(3)).
- a. ☐ are transmitted herewith (required only if not transmitted the International Bureau)
 - b. ☐ have been transmitted by the International Bureau.
6. ☐ A translation of the amendments to the claims under PCT Article 19 (35 USC 371(c)(3)).
7. ☒ An oath or declaration of the inventor (35 USC 371(c)(4)).
8. ☐ A translation of the annexes to the International Preliminary Examination Report under PCT Article 36 (35 USC 371(c)(5)).

Other document(s) or information included:

- 9. ☐ An information Disclosure Statement under 37 CFR 1.97 and 1.98.
- 10. ☒ An assignment document for recording.
- 10.1 ☐ A copy of the International Filing Receipt.
- 10.2 ☒ A copy of the International Search Report.
- 10.3 ☒ A copy of the International Preliminary Examination Report.
- 10.4 ☒ Preliminary Amendment deleting multiple dependent claims.
- 11. The above checked items are being transmitted:
 - ☒ by thirty (30) months and a proper demand for International Preliminary Examination was made by the 19th month from the earliest claimed priority date.
- 12. At the time of transmittal, the time limit for amending claims under Article 19:
 - a. ☐ has expired and no amendments were made.
 - b. ☐ has not yet expired.
- 13. Certain requirements under 35 USC 371 were previously submitted by the applicant on _____, namely:

Susan S. Jackson

Susan S. Jackson/Registration No. 41,302
 Clariant Corporation
 4331 Chesapeake Drive
 Charlotte, North Carolina 28216
 Telephone: 704/395-6716
 Facsimile: 704/395-6727

Date: December 10, 1998

0966671 060301
 106990 1/989960

Attorney's Docket No. 153-5916/PCT

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re new U.S. patent application of : Attn: BOX PCT (DO/EO/US)
Jan MALIK et al. :
Serial No. To Be Assigned :
Filed: December 10, 1998 :
For: STABILIZER COMPOSITIONS : December 10, 1998

PRELIMINARY AMENDMENT

Assistant Commissioner for Patents
Washington, DC 20231

Dear Sir:

Preliminary to the examination of the above-identified application, please
amend the application as follows:

In the Specification:

Page 1, delete lines 4-5.

Page 1, line 6, insert --BACKGROUND OF THE INVENTION--.

Page 2, line 1, insert --SUMMARY OF THE INVENTION--.

Page 2, line 2, after "SUMMARY OF THE INVENTION", insert --The invention
relates to stabilizer compositions for stabilizing polyethylene-based thermoplastic
polymers in processing.--

Page 2, line 23, insert --DESCRIPTION OF THE PREFERRED EMBODIMENTS--.

In the Claims:

Please amend the claims as follows:

Claim 3, line 1, delete "or 2".

Claim 4, line 1, delete "any preceding claim" and substitute therefor --claim
1--.

Claim 5, line 1, delete "any preceding claim" and substitute therefor --claim
1--.

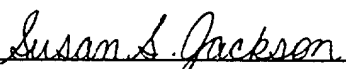
Claim 8, line 2, delete "or".

In re new U.S. patent application of:
Jan MALIK et al.
Serial No. To Be Assigned
Filed: December 10, 1998
Page 2

REMARKS

The claims have been amended to delete the multiple dependent claims so
as to reduce the application fees.

Respectfully submitted,



Susan S. Jackson
Registration No. 41,302

Clariant Corporation
Industrial Property Department
4331 Chesapeake Drive
Charlotte, North Carolina 28216
Telephone: 704/395-6716
Facsimile: 704/395-6727

09868871:060501
T05090:12888860

STABILIZER COMPOSITIONS

5 The invention relates to stabilizer compositions for stabilizing polyethylene-based thermoplastic polymers in processing.

During processing into articles, polymers are subjected to high temperatures and pressures which can have an adverse effect on the molecular weight and physical properties of the polymers and on the appearance of the finished articles made from these polymers. Therefore, there is a need for antioxidant and color stabilizing additives which minimize oxidation and discoloration.

United States Patent No. 4 806 580 discloses stabilizer mixtures for stabilizing plastics material, e.g. polypropylene, in processing, containing (a) a chroman derivative and (b) an organic phosphite or phosphonite in a weight ratio of a:b of 1:5 to 1:14.

European Patent Application 0 542 108 discloses a stabilizing mixture for plastics materials containing polyethylene, comprising (a) a tocopherol compound and (b) a phosphorus- or sulphur-containing secondary antioxidant in a weight ratio of a:b of 1:1.4 to 1:5. The corresponding disclosure exists in the article of S.F. Laermer and P.F. Zambetti published in the
20 Journal of Plastic Film & Sheeting, volume 8, 1992, pages 228 to 248. This article further describes a mixture of α -tocopherol, "Irganox (trade mark) 1010" (a sterically hindered phenol), and "Irgafos 168" (a phosphite) in the weight ratio of 1:1:4 for the stabilization of polypropylene.

The article of S.S. Young, S.F. Laermer and P.F. Zambetti published in the Journal of Plastic
25 Film & Sheeting, volume 11, 1995, pages 126 to 142, demonstrates that α -tocopherol formulations can replace phenol/phosphite combinations as the primary antioxidant system, or the α -tocopherol formulations can simply replace the phosphite portion to achieve better performance.

30 German Patent No. 26 60 746 discloses the use of symmetrical triarylphosphites in combination with sterically hindered phenolic antioxidants in the stabilization of polyolefins.

It is the object of the present invention to provide an improved stabilizer composition for the stabilization of polyethylene-based thermoplastic polymers, such that both processing stability and color of the final products are improved.

5 This object is achieved with the stabilizer composition as hereinunder described. It has surprisingly been found that relatively small additions of α -tocopherol to known additive systems for polyethylene-based polymers, composed of a phenolic antioxidant and a phosphorus based secondary antioxidant, results in an unexpected synergistic effect. This stabilizer composition according to the invention comprising at least one member of three known additive classes significantly outperforms the additive systems that are known in the art.

Accordingly, the present invention provides a stabilizer composition comprising

- 10
15
20
25
30
- a) at least one sterically hindered phenol,
 - b) at least one phosphorus-containing secondary antioxidant, and
 - c) at least one tocopherol compound
- wherein the weight ratio of component (a) to component (b) is from 2:1 to 1:4 and the weight ratio of component (a) to component (c) is from 2:1 to 10:1,

for the stabilization of polyethylene-based thermoplastic polymers against degradation, crosslinking and/or discoloration due to the exposure to heat or light, especially in the presence of oxygen.

The preferred weight ratio of component (a) to component (b) is 1:1 and that of component (a) to component (c) is 5:1.

The term "tocopherol compound" refers to any compound having the basic tocopherol structure of the vitamin E group. The preferred tocopherol compound as component (a) in the stabilizer composition according to the present invention is α -tocopherol (5,7,8-trimethyl-tocol).

As used herein, the term "sterically hindered phenol" refers to those compounds that are preferably derived from 2,6-di-tert.-butyl-phenol, 2-tert.-butyl-6-methylphenol, 2-tert.-butyl-5-

methylphenol or other hindered phenols. Preferred examples of such compounds are 2,2'-Bis[3,5-[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]-1-oxopropoxy]methyl-1,3-propanediyl-3,5-bis(1,1-dimethylethyl)-4-hydroxybenzenepropanoate; octadecyl-3,5-bis(1,1-dimethylethyl)-4-hydroxybenzene-propanoate; 1,3,5-tris[[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]methyl]-1,3,5-triazine-2,4,6(1H,3H,5H)-trione; 4,4',4''-[2,4,6-trimethyl-1,3,5-benzenetriyl]tris-(methylene)]tris[2,6-bis(1,1-dimethylethyl)-phenol]; Ethanediyl-3,5-bis(1,1-dimethylethyl)-4-hydroxy-thiodi-2,1-benzenepropanoate; 2:1 calcium salt of monoethyl-[[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]methyl]-phosphonic acid ester; 2-[3-[3,5-bis(1,1-dimethylethyl)-4-hydroxy-phenyl]-1-oxo-propyl]hydrazide-3,5-bis(1,1-dimethylethyl)-4-hydroxy-benzenepropanoic acid; 2,2'-oxamido-bis-[ethyl-3-(3,5-di-tert.butyl-4-hydroxyphenyl)propionate] or mixtures thereof.

As also used herein, the term "phosphorus-containing secondary antioxidant" refers to compounds being hydroperoxide decomposers, i.e. compounds having the ability to react with hydroperoxides to yield non-radical products, essentially decomposing hydroperoxides into stable by-products. Examples of such compounds are triesters of phosphorous acid (phosphites) and diesters of phosphorous acid (phosphonites). Preferred specific members of the phosphorus-containing secondary antioxidants are Triphenylphosphite, Trisisodecylphosphite; Tris(nonylphenyl)phosphite; Distearyl pentaerythritol diphosphite; 2,4,6-tri-tert.-butyl-phenyl-2-butyl-2-ethyl-1,3-propanediol phosphite; Bis(2,4-di-tert.-butylphenyl)-pentaerythrityl diphosphite; 2,2',2''-nitriolo triethyl-tris[3,3',5,5'-tetra-tert.-butyl-1,1'-biphenyl-2,2'-diyl]phosphite; Bis[2,4-di-tert.-butyl-6-methyl-phenyl]ethyl phosphite; 2,2'-Ethylidene-bis-(4,6-di-tert.-butylphenyl)fluorophosphite; Tris-(2,4-di-tert.-butylphenyl)phosphite; the 4,6-di-tert.-butyl-m-cresol condensation products with the Friedel-Crafts-reaction products of biphenyl and phosphorus trichloride; Tetrakis [2,4-di-tert.-butylphenyl]-4,4'-biphenylenediphosphonite; the condensation products of 2,4-di-tert.-butylphenol with the Friedel-Crafts-reaction product of biphenyl and PCl_3 .

The term "polyethylene-based thermoplastic polymer" refers to all types of homo- and copolymers of ethylene, e.g. high density polyethylene, low density polyethylene, linear low density polyethylene, ultra low density polyethylene and ultra high molecular weight polyethylene or to blends of polyethylene and at least another polymer. Those skilled in the art

will know how to make the different types of copolymers that can be formed with ethylene, including alternating, block and graft copolymers as well as how to make blends of polyethylene with other polymers.

- 5 If in the stabilizer composition according to the present invention more than one compound of component (a), (b) or (c) is present, the definition of the weight ratios always refers to the total amount of component (a), (b) or (c).

The phosphites and phosphonites as well as the sterically hindered phenols disclosed above can be synthesized by techniques well known in the art from known compounds. They are commercially available and their tradenames as well as their chemical structural formulae are summarized in table 1.

The stabilizer composition of the present invention may be produced by simple physical mixing of the compounds (a), (b) and (c) in the desired weight ratio according to known methods. A preferred method is the dry blending of the components resulting in a free-flowing mixture which may be further subjected to a pre-extrusion step in order to achieve homogeneity and easy-to-handle pellets. The process for producing the stabilizer composition is a further aspect of the present invention.

20

A masterbatch composition comprising a stabilizer composition according to the invention and a thermoplastic material which is identical to or compatible with the thermoplastic polymer to be stabilized is similarly an object of the invention. Masterbatch compositions according to the invention comprise 10 to 80% by weight, preferably 5 to 25% by weight of the stabilizer
25 composition and 90% to 20% by weight, preferably 95 to 75% by weight of the afore-mentioned thermoplastic material.

30

The invention also provides a method for enhancing the processing stability of polyethylene-based thermoplastic polymers comprising - incorporating therein at any convenient state of the processing - a stabilizing quantity of the stabilizer composition. This can be carried out according to known methods and may involve the incorporation of the stabilizer composition as such or in

the form of the above-mentioned masterbatch. Also, the individual components can be incorporated separately in the correct ratio.

The concentration of the stabilizer mixture amounts from 0.001 to 5% , preferably from 0.01 to 1%, more preferably from 0.1 to 0.5% by weight, based on the weight of the polyethylene-based thermoplastic polymers to be stabilized.

The present invention further embraces a stabilized polyethylene-based thermoplastic polymer and any article manufactured therefrom that has been stabilized by the incorporation therein a stabilizing quantity of the stabilizer composition according to the present invention.

Further additives which may be added, if appropriate, include plasticizers, lubricants, emulsifiers, pigments, dyestuffs, nucleating agents, optical brighteners, flameproofing agents, antistatic agents, blowing agents, U.V. absorbers, U.V. quenchers, hindered amine light stabilizers, metal deactivators and several others commonly used.

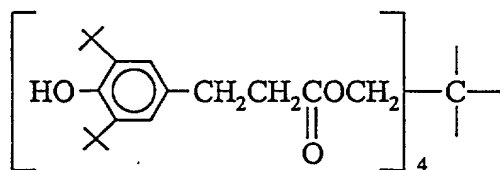
As mentioned above the tradenames, the chemical structural formulae and the chemical names of the components (a) and (b) are listed in table 1.

TABLE 1

Trade names, chemical names and chemical structural formulae of the individual components of the stabilizer composition according to the invention.

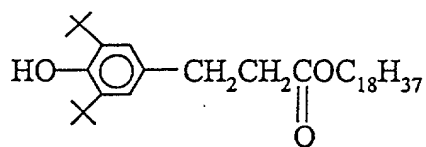
Irganox[®] 1010 (Ciba-Geigy)

Tetrakis[methylen-3-(3',5')-di-tert.-butyl-4'-hydroxyphenyl]propionate]methane



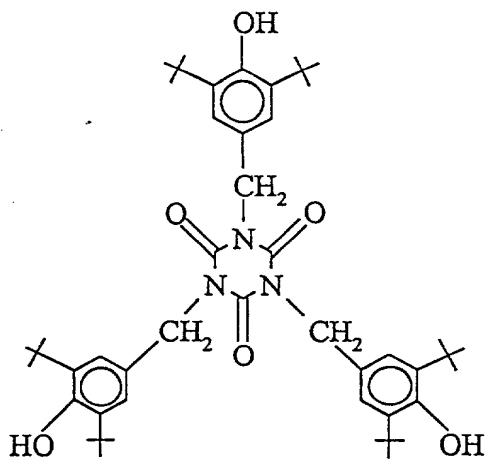
Irganox®1076 (Ciba-Geigy)

Octadecyl-3,5-bis(1,1-dimethylethyl)-4-hydroxybenzenepropanoate



Irganox®3114 (Ciba-Geigy)

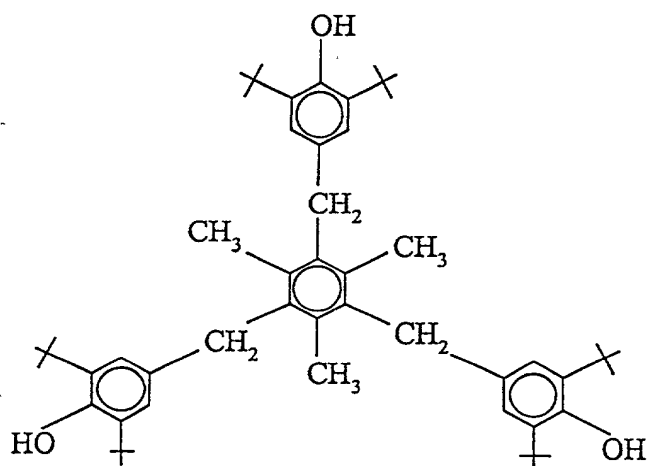
1,3,5-tris[[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]methyl]-1,3,5-triazine-2,4,6(1H,3H,5H)-trione



Irganox® 1330 (Ciba-Geigy)

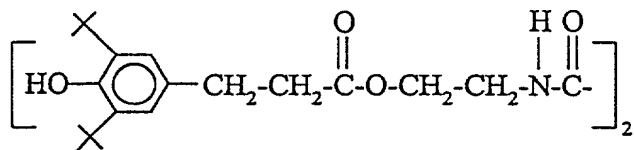
4,4',4''-[(2,4,6-trimethyl-1,3,5-benzenetriyl)tris(methylene)]-tris[2,6-bis(1,1-dimethylethyl)phenol]

5



Naugard® XL-1 (Uniroyal Chemical)

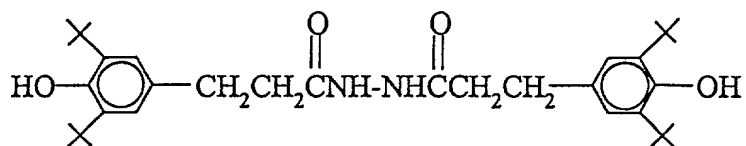
2,2'-oxamido-bis-[ethyl-3-(3,5-di-tert.-butyl-4-hydroxyphenyl)propionate]



Irganox® MD-1024 (Ciba-Geigy)

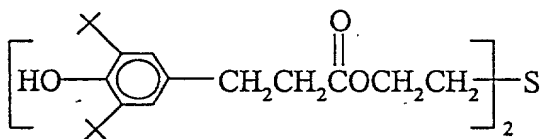
2-[3-[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]-1-oxpropyl]hydrazide-3,5-bis(1,1-dimethylethyl)-4-hydroxy-benzenepropanoic acid

5



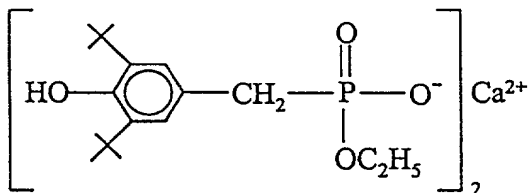
Irganox® 1035 (Ciba-Geigy)

Ethanediyyl -3,5-bis(1,1-dimethylethyl)-4-hydroxy-thiodi-2,1-benzenepropanoate



15 Irganox® 1425 WL (Ciba-Geigy)

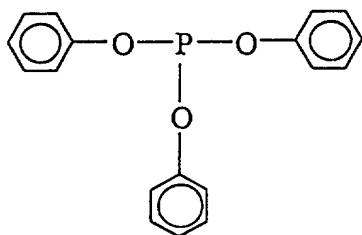
2:1 calcium salt of monoethyl-[[3,5-bis(1,1-dimethylethyl)-4-hydroxy-phenyl]methyl]-phosphonic acid ester



20

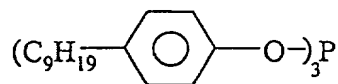
Weston® TPP (General Electric)

Triphenyl phosphite



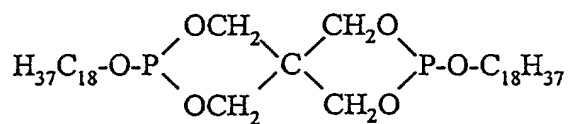
Weston® TNPP (General Electric)

Tris(nonylphenyl)phosphite



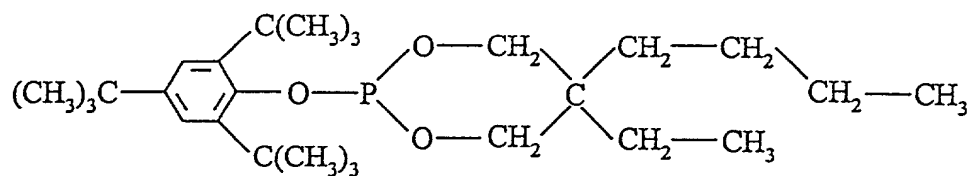
Weston® 618 (General Electric)

15 Distearyl pentaerythritol Diphosphite



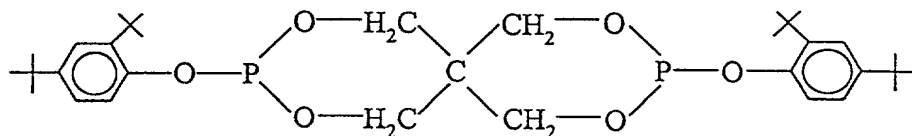
Ultranox® 641 (General Electric)

2,4,6-tri-tert.-butylphenyl-2-butyl-2-ethyl-1,3-propanediol phosphite



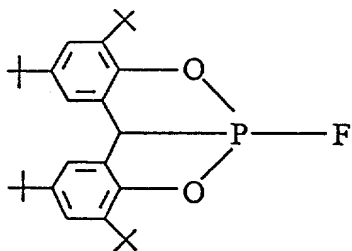
Ultranox® 626 (General Electric)

Bis(2,4-di-tert-butylphenyl)-pentaerythritol-diphosphite



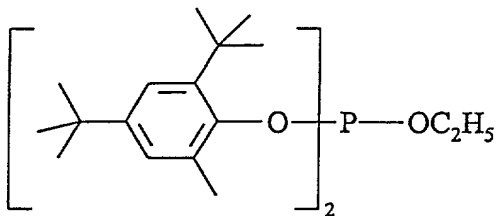
Ethanox 398 (Ethyl Corporation)

2,2'-Ethylidene-bis(4,6-di-tert.-butylphenyl)fluorophosphite



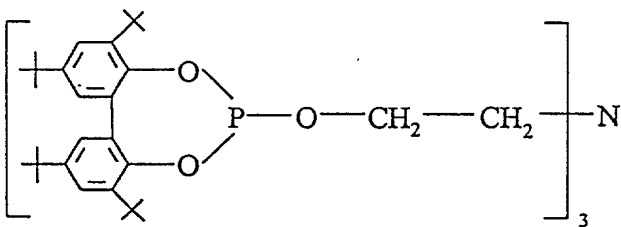
Irgafos®38 (Ciba-Geigy)

Bis[2,4-di-tert.-butyl-6-methylphenyl]ethyl phosphite



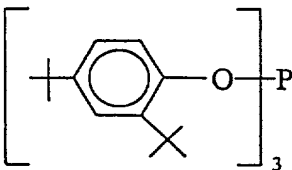
Irgafos®12 (Ciba-Geigy)

2,2',2''-nitrilo triethyl-tris[3,3',5,5'-tetra-tert.-butyl-1,1'-biphenyl-2,2'-diyl]phosphite



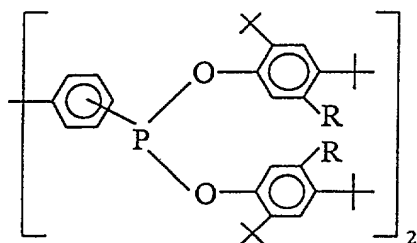
Irgafos®168 (Ciba-Geigy)

15 Tris(2,4-di-tert.-butylphenyl)phosphite



GSY P-101 (Yoshitomi)

Tetrakis[2,4-di-tert.-butyl-5-methyl-phenyl]biphenylene-diphosphonite

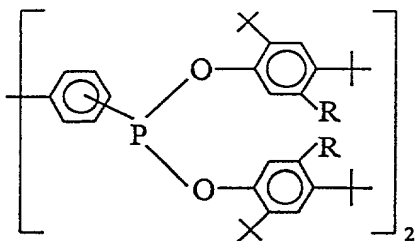
R = CH₃

Irgafos® P-EPQ (Ciba-Geigy)

Sandostab® P-EPQ (Clariant)

Composed of

- 1) 50-80 parts of Tetrakis(2,4-di-tert.-butylphenyl)-biphenylene-diphosphonite



R = H

- 2) 10-25 parts of bis(2,4-di-tert.-butylphenyl)biphenylene-monophosphonite
- 3) 10-25 parts of tris(2,4-di-tert.-butylphenyl)phosphite

For avoidance of doubt, in this specification tert.-butyl means tertiary-butyl, -C(CH₃)₃.

- The following non-limiting examples illustrate the invention in its various aspects. All parts and percentages are expressed by weight.

EXAMPLES

The following types of unstabilized polyethylene are used in the examples described hereinunder:

PE No.	Type of Polyethylene
PE-1	HDPE, prepared with Cr-catalyst, blow molding grade MFI = 0.25 g / 10 min (190°C / 2.16 kg) MFI = 18.96 g / 10 min (190°C / 21.6 kg)
PE-2	HDPE, prepared with Ti-catalyst MFI = 2.4 g / 10 min (190°C / 10 kg)
PE-3	LLDPE, MFI = 1.4 g / 10 min (190°C / 2.16 kg)

The various additives tested are used as received from the respective suppliers. ATP means α -tocopherol.

Code	Composition component	Tradename / Manufacturer
AO-1	component (a)	Irganox 1010/Ciba-Geigy
AO-2	component (a)	Irganox 1076/Ciba-Geigy
PS-1	component (b)	Sandostab P-EPQ/Clariant
ATP	component (c)	Ronotec 201/Hoffmann-La Roche

- 10 The tested samples are prepared as follows. In each case 100 parts of the different types of polyethylenes are dry-blended with the tested additives as well as with a lubricant, followed by pre-extrusion in a single screw laboratory extruder with 80 rpm at temperatures as given below in table 2.
- 15 After extrusion compounding the materials are sequentially passed in total five times through a single screw extruder (compression ratio = 1:3, L/D = 20) at temperatures and numbers of revolutions per minute as given below in table 2. The unstabilized resins are in most cases also processed through the extruder so that all materials have the same heat history. After the five

extrusion passes the materials are assessed in respect of color quality and change of molecular weight.

The color quality is reported in terms of the Yellowness Index (YI), determined on the granules in accordance with the ASTM 1925-70 Yellowness Test. The higher the value, the lower the color quality, i.e. the worse the discoloration.

The change of the molecular weight is reported in terms of the melt flow index (MFI) according to ASTM D-1238-70, measured in a Zwick melt flow indexer at temperatures and loads as given below. Experiment errors of the MFI measurements are determined from an independent series of measurements, and are found to be ± 0.01 for the MFI measured at 190°C and 2.16 kg load, and ± 0.3 for the MFI measured at 190°C and 21.6 kg load.

Comparison data refers to the following prior art documents (D):

Code	Document
D-1	Gächter, Müller, Plastics Additives, 3rd edition, Hanser Publishers 1990
D-2	Journal of Plastic Film & Sheeting, volume 11, 1995, pages 126-142
D-3	US P 4,806,580
D-4	EP-A-0 542 108

TABLE 2

Additional conditions of the sample preparations.

Example No.	Type of PE	Lubricant	Compounding temperature	Extrusion	
				temp.	rpm
1	PE-1	500 ppm Zn Stearate	200°C	220°C	70
2	PE-1	500 ppm Zn Stearate	200°C	240°C	70
3	PE-2	1000 ppm Ca Stearate	210°C	270°C	100
4	PE-3	1000 ppm Ca Stearate	210°C	240°C	70

EXAMPLE 1

Comparison data (CD) CD-1 presents the stabilization system according to the currently used formulation.

CD-2 presents the system described in D-2.

CD-3 corresponds to the disclosures given in D-3 and D-4.

I-1 is an example according to the present invention.

As is evident from the results given in table 3, the stabilizer composition according to the invention offers significantly better melt stabilization than systems according to the prior art, providing simultaneously an excellent color retention. The unstabilized polymer (PE-1) illustrates the effect of pronounced crosslinking after 5 repeated extrusions; the MFI drops to the undesirable value of 9.96 g / 10 min.

TABLE 3

Data of example 1.

Experiment	AO-1	PS-1	ATP	MFI		YI
	[ppm]	[ppm]	[ppm]	[190°C/2.16 kg]	[190°C/2.16 kg]	
PE-1	---	---	---	0.05 g / 10 min	9.96 g / 10 min	1.23
CD-1	500	500	---	0.16 g / 10 min	15.79 g / 10 min	-0.6
CD-2	500	---	100	0.15 g / 10 min	15.79 g / 10 min	0.57
CD-3	---	500	100	0.11 g / 10 min	13.06 g / 10 min	0.95
I-1	500	500	100	0.2 g / 10 min	17.43 g / 10 min	-0.55

EXAMPLE 2

Comparison data CD-1 to CD-4 describe common formulations of polyethylene.

CD-5 and CD-6 correspond to the additive systems described in D-2.

CD-7 and CD-8 correspond to the systems disclosed in D-3.

I-1 to I-4 are stabilizer compositions according to the invention.

As can be seen from the results summarized in table 4, the unstabilized polymer is significantly crosslinked after the 5 extrusions. Further, the stabilizer compositions according to the invention significantly outperform the systems known in the art with regard to the melt flow retention and color stabilization. The comparison of the results of CD-3 with those of I-1 on the one hand and of the results of CD-4 with those of I-2 on the other hand clearly demonstrate that the obtained stabilization effect is not caused by an increase of the total concentration of phenolic antioxidant (AO-1 and ATP). It is believed that the stabilization performance of the stabilizer compositions according to the invention, i.e. I-1 and I-2 in the aforementioned comparison, is achieved by a synergistic action of all three components of the stabilizer composition and, thus, cannot be regarded as pure additive, i.e. quantity depending effects. Since the effectiveness of the single components AO-1, PS-1 and ATP is well known in the prior art, it is surprising to find this particular synergy by triple-combinations according to the present invention.

TABLE 4

Data of example 2.

Experiment	AO-1	PS-1	ATP	MFI		YI
	[ppm]	[ppm]	[ppm]	[190°C/2.16 kg]	[190°C/2.16 kg]	
PE-1	---	---	---	0.01 g / 10 min	9.10 g / 10 min	---
CD-1	500	750	---	0.17 g / 10 min	16.76 g / 10 min	-1.00
CD-2	500	1000	---	0.20 g / 10 min	18.03 g / 10 min	-1.10
CD-3	750	750	---	0.17 g / 10 min	17.01 g / 10 min	-0.11
CD-4	750	1000	---	0.20 g / 10 min	18.21 g / 10 min	-0.24
CD-5	500	---	100	0.14 g / 10 min	15.70 g / 10 min	-0.39
CD-6	750	---	100	0.15 g / 10 min	16.33 g / 10 min	-1.14
CD-7	---	750	100	0.14 g / 10 min	14.49 g / 10 min	0.59
CD-8	---	1000	100	0.16 g / 10 min	15.46 g / 10 min	0.42
I-1	500	750	100	0.21 g / 10 min	18.16 g / 10 min	-0.90
I-2	500	1000	100	0.23 g / 10 min	18.98 g / 10 min	-1.10
I-3	750	750	100	0.22 g / 10 min	18.55 g / 10 min	-1.61
I-4	750	1000	100	0.24 g / 10 min	19.10 g / 10 min	-1.67

5

EXAMPLE 3

CD-1 and CD-2 are comparison tests. I-1 is according to the invention. The results are summarized in table 5.

10

Again, the stabilizer composition according to the invention gives the best melt and color stabilization. The comparison of the results of CD-2 or CD-3 with those of I-1 repeatedly appears to confirm that the stabilization effect is achieved by a synergistic effect of all three components of the stabilizer composition according to the invention.

15

TABLE 5

Data of example 3.

Experiment	AO-1 [ppm]	PS-1 [ppm]	ATP [ppm]	MFI [190°C/10 kg]	YI
PE-2	---	---	---	4.83 g / 10 min	---
CD-1	1000	---	---	3.97 g / 10 min	2.8
CD-2	500	1000	---	2.86 g / 10 min	2.7
I-1	400	1000	100	2.55 g / 10 min	1.4

EXAMPLE 4

CD-1 and CD-2 are comparison tests. I-1 is according to the invention. The results are summarized in table 6.

As can be seen from table 6 the stabilizer composition according to the invention conferred unexpectedly outstanding melt flow retention and color stabilization to the tested polymer. This stabilization effect is achieved at a significantly lower total stabilizer concentration than it was used in CD-2.

TABLE 6

Data of example 4.

Experiment	AO-2 [ppm]	PS-1 [ppm]	ATP [ppm]	MFI [190°C/2.16 kg]	YI
CD-1	700	---	---	0.82 g / 10 min	3.0
CD-2	700	1000	---	1.25 g / 10 min	0.2
I-1	250	1000	100	1.30 g / 10 min	-0.8

CLAIMS

1. A stabilizer composition for the stabilization of polyethylene-based thermoplastic polymers comprising
- a) at least one sterically hindered phenol,
- b) at least one phosphorus-containing secondary antioxidant, and
- c) at least one tocopherol compound
- wherein the weight ratio of component (a) to component (b) is from 2:1 to 1:4 and the weight ratio of component (a) to component (c) is from 2:1 to 10:1.
2. A composition according to claim 1 wherein the weight ratio of component (a) to component (b) is 1:1 and the weight ratio of component (a) to component (c) is 5:1.
3. A composition according to claim 1 or 2 wherein the tocopherol compound is α -tocopherol (5,7,8-Trimethyl-tocol).
4. A composition according to any preceding claim wherein the sterically hindered phenol is 2,2'-Bis[3,5-[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]-1-oxoprop-oxy]methyl-1,3-propanediyl-3,5-bis(1,1-dimethylethyl)-4-hydroxybenzenepro-panoate; Octadecyl-3,5-bis(1,1-dimethylethyl)-4-hydroxybenzenepropanoate; 1,3,5-tris[[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]methyl]1,3,5-triazine-2,4,6(1H,3H,5H)trione; 4,4',4''-[2,4,6-trimethyl-1,3,5-benzenetriyl]tris-(methylene)]tris[2,6-bis(1,1-dimethylethyl)-phenol; Ethanediyl-3,5-bis(1,1-dimethylethyl)-4-hydroxy-thiodi-2,1-benzenepropanoate; 2:1 calcium salt of monoethyl-[[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]-methyl]-phosphonic acid ester; 2-[3-[3,5-bis(1,1-dimethylethyl)-4-hydroxy-phenyl]-1-oxopropyl]-hydrazide-3,5-bis(1,1-dimethylethyl)-4-hydroxy-benzene-propanoic acid; 2,2'-oxamido-bis-[ethyl-3-(3,5-di-tert.-butyl-4-hydroxyphenyl)-propionate] or mixtures thereof.

5. A composition according to any preceding claim wherein the phosphorus-containing secondary antioxidant is Triphenylphosphite, Tris-isodecylphosphite; Tris(nonylphenyl)phosphite; Distearyl pentaerythritol diphosphite; 2,4,6-tri-tert.-butylphenyl-2-butyl-2-ethyl-1,3-propanediol phosphite; Bis(2,4-di-tert.-butylphenyl)-pentaerythrityl diphosphite; 2,2',2''-nitrilo triethyl-tris[3,3',5,5'-tetra-tert.-butyl-1,1'-biphenyl-2,2'-diyl]phosphite; Bis[2,4-di-tert.-butyl-6-methyl-phenyl]ethyl phosphite; 2,2'-Ethylidene-bis-(4,6-di-tert.-butylphenyl)fluorophosphite; Tris(2,4-di-tert.-butylphenyl)phosphite; the 4,6-di-tert.-butyl-m-cresol condensation products with the Friedel-Crafts-reaction products of biphenyl and phosphorus trichloride; Tetrakis [2,4-di-tert.-butylphenyl]-4,4'-biphenylenediphosphonite; the condensation products of 2,4-di-tert.-butylphenol with the Friedel-Crafts-reaction product of biphenyl and PCl_3 .
6. A method for enhancing the processing stability of polyethylene-based thermoplastic polymers comprising incorporating therein before or during processing a stabilizing quantity of the stabilizer composition according to claim 1.
7. A method according to claim 6 wherein the stabilizer composition is added in an amount of from 0.001 to 5% by weight, preferably from 0.01 to 1% by weight, more preferably from 0.1 to 0.5% by weight, based on the thermoplastic polymer.
8. A process for producing a stabilizer composition according to claim 1 comprising mixing the components (a), (b) and (c) in the weight ratios given in claim 1 or
9. A masterbatch composition comprising a stabilizer composition according to claim 1 and a thermoplastic material which is identical or compatible with the polyethylene-based thermoplastic polymer to be stabilized.

10. A masterbatch composition according to claim 9 comprising 10 to 80% by weight, preferably 5 to 25% by weight of the stabilizer composition and 90 to 20% by weight, preferably 95 to 75% by weight of said thermoplastic material.

5

11. A stabilized polyethylene-based thermoplastic polymer and any article manufactured therefrom being stabilized by the incorporation therein of a stabilizing quantity of the stabilizing composition according to claim 1.

10

0986871 060501
103090 1 689860

**DECLARATION AND POWER OF ATTORNEY FOR
UNITED STATES PATENT APPLICATION**

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name, and

I believe that I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if more than one name is listed below) of the subject matter which is claimed and for which a United States patent is sought on the invention entitled

STABILIZER COMPOSITIONS

the specification of which

☒ is attached hereto.

☐ was filed on as application and, if these brackets contain an X ☐, was amended on

☒ was filed as Patent Cooperation Treaty international application under No: **PCT/IB97/007700 on 24 June 1997**, if these brackets contain an X ☐, was amended under Patent Cooperation Treaty Article 19 on , if these brackets contain an X ☐, entered the national stage in the United States on , 19 and was accorded Serial No. , and, if these brackets contain an X ☐, was amended on

I hereby state that I have reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment(s) referred to above.

I acknowledge my duty to disclose all information which is known by me to be material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I hereby claim the benefit under Title 35, United States Code, §119(a)-(d) of any foreign application(s) for patent or inventor's certificate indicated below and of any Patent Cooperation Treaty international application(s) designating at least one country other than the United States indicated below and have also identified below any foreign application(s) for patent or inventor's certificate and Patent Cooperation Treaty international application(s) designating at least one country other than the United States for the same subject matter and having a filing date before that of the application for said subject matter the priority of which is claimed:

09863871.060301

<u>Country</u>	<u>Number</u>	<u>Filing Date</u>	<u>Priority Claimed</u>
GB	9613515.7	27. June 1996	[X]Yes [] No

I hereby claim the benefit under Title 35, United States Code, §119(e) of any United States provisional application(s) listed below:

Application

Serial No.

Filed

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and of any Patent Cooperation Treaty international application(s) designating the United States listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in said prior application(s) in the manner required by the first paragraph of Title 35, United States Code, §112, I acknowledge my duty to disclose all information known to me to be material to patentability as defined in Title 37, Code of Federal Regulations, §1.56 which became available between the filing date(s) of the prior application(s) and the national or Patent Cooperation Treaty international filing date of this application:

Application

Serial No.

Filed

Status (Pending, Abandoned or Patented) _____

I hereby appoint the following:

PHILIP P. MCCANN

Reg. No. 30,919

THOMAS C. DOYLE

Reg. No. 22,340

GABRIEL LOPEZ

Reg. No. 28,440

HESNA J. PFEIFFER

Reg. No. 22,640

MILES B. DEARTH

Reg. No. 35,115

SCOTT E. HANF

Reg. No. 38,906

I hereby appoint the following (cont.):

SUSAN JACKSON

Reg. No. 41,302

ANDREW F. SAYKO, Jr.

Reg. No. 22,827

respectively and individually, as my attorneys and/or agents, with full power of substitution and revocation, to prosecute this application and to transact all business in the Patent and Trade-mark Office connected therewith. Please address all communications to:

PHILIP P. MCCANN, CLARIANT CORPORATION, 77 Center Drive, Building 5200,
Charlotte, NC28217, whose telephone number is 704-559 6121.

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

IMPORTANT: Before this declaration is signed, the patent application (the specification, the claims and this declaration) must be read and understood by each person signing it, and no changes may be made in the application after this declaration has been signed.

Sole inventor or

first joint inventor:

Full name : Jan MALIK

Signature :

Date :

Citizenship :

Residence :

P.O. Address :

Jan MALIK

Jan MALIK

16th day of September 1998

Slovak Republic

17, rue Charles Peguy, F-68300 St. Louis, France

Same as above

FR

Second joint inventor,

if any:

2-00

Full name : Klaus STOLL

Signature :

Klaus Stoll

Date :

16th day of September, 1998

Citizenship :

German

Residence :

Am Sonnenrain 18. D-79589 Binzen, Germany

P.O. Address :

Same as above

DE

Third joint inventor,

if any:

3-00

Full name : Andreas THUERMER

Signature :

Andreas Thuermer

Date :

16th day of September 1998

Citizenship :

German

Residence :

2, Rue Chancel, F-68330 Huningue, France

P.O. Address :

Same as above

FR

Fourth joint inventor,

if any:

Full name :

Signature :

Date :

Citizenship :

Residence :

P.O. Address :

Same as above

Fifth joint inventor,

if any:

Full name :

Signature :

Date :

Citizenship :

Residence :

P.O. Address :

T05090 T/089860